92-018148/03 D25 E14

1NSF 09.07.90 *EP -466-558-A D(!1-A) E(10-J2B1) N(6-A)

INST FRANCAIS DU PETROLE

09.07.90-FR-008801 (15.01.92) C07c-02/66 C07c-15/19

2- and 3-Phenyl alkane(s) prepn. - by reacting benzene with linear olefin using catalyst of de-aluminised mordenite, for biodegradable detergent

C92-007844 RIAT BE DE ES GB IT NU

Prepn. of a mixt. of 2-phenylalkanes and 3-phenylalkane comprises reacting, in a reaction zone, benzene with a charge contg. at least one 9-16C linear olefin using a catalyst based on dealuminated mordenite (I).

The obtd. prod. is then fractionated to obtain separate-

ly.
n first fraction contg. unreacted benzene;
n second fraction contg. unreacted 9-16C olefin;
n third fraction contg. a mixt. of the 2- and 3-phenylal-kanes; and

a fourth fraction contg. at least one poly-alkylbenzene.

The fourth fraction is at least partly recycled to the reaction zone where it reacts with benzene in presence of (1) to be at least partly transalkylated, and a mixt. of 2-and 3-phenylalkanes is recovered.

(I) has

overall Si/Al atomic ratio of 15-85(pref. 15-60).
No wt. content of <1000 (pref. <250) ppm
a very low or zero content of extra-network Al species:
an elementary mesh vol. <2.760 (pref. <2.745) nm³.
asymmetrical vibration frequency of the tetrahedra TO₄
(T = Al or Si) >1070 (pref. >1082) cm¹
in the IR spectrum; and
micropore vol. > 0.150 (pref. > 0.190) cm³ (liquic)/g.

The first and second fractions are pref. also at least partially recycled. Reaction is pref. at <60°C.

USE

The 2- and 3-phenyl alkane mixt. is used as base for prodn. of biodegradable detergents (by sulphonation).

ADVANTAGE

Catalytic performance is superior to prior art catalysts; (I) is highly active, highly selective and esp. resistant to deactivation.

PREFERRED CATALYST

(1) is such that the intensity of signals corresp. to the outside frame (extra network) alumina species FP-466558-A+

to the intensity of signals corresp. to the frame alumina species (measured by 27 Al, NMR and IR spectroscopy in the OH gps. region) is <0.15 (pref. <0.05).

(1) has a secondary pore network of above 0.020 cm³/g. The catalyst pref. also contains a matrix selected from clays, alumina, silica, magnesia, zirconia, TiO₂ and/or boron oxide.

EXAMPLE

100g of a large pore mordenite in Na form (TSZ 600 NAA from Tosoh Cc.) of formula Na₂AlO₂(SiO₂)_{5·1} was heated at reflux at 100°C for 2 hrs. in an ammonium nitrate soln.

(4M), the vol. of which was four times the wt. of mordenite.

This cation exchange operation was repeated three times. The prod. (contg. 500 ppm Na) was treated with an uq. soln. of HNO₃ of normality 4.5N at reflux. The obtd. catalyst had the following characteristics. Na content:- 25 ppm; Si/Al (overall):- 40; elementary mesh vol.:-2.715 nm³; IR frequency TO₄:- 1086 cm⁻¹; micropore vol.:- 0.272 cm³/g and mesopore vol.:- 0.178 cm³/g (Dubinin).

The catalyst was used for the alkylation of benzene with dodecane-1 under the following conditions:- temp. 50°C; pressure 4MPa, LHSV 3, benzene/dodecene-1 molar ratio: 5.5.

Selectivities to 2-phenyl-dodecane and 3-phenyldodecane w.r.t. converted dodecene-1 were 75.8% and 10.6% resp. (14pp950DAHDwgNo0/1).

(F) ISR: US4731497 EP-366515 FR2084704

EP-466558-A